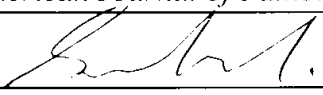


FORM PTO-1449 (REV. 7-80)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTY. DOCKET NO. 240083.501D6		EXPRESS MAIL NO. EL615486917US	
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				FILING DATE October 25, 2000		GROUP ART UNIT 1636	
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	AA						
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SE	AB	WO 98/45435	10/15/98	PCT			
	AC						
	AD						
	AE						
	AF						
OTHER PRIOR ART (Including Author, Title, Date, Pertinent Pages, Etc.)							
SE	AG	Barbosa et al., "Identification of the Homologous Beige and Chediak-Higashi Syndrome Genes," <i>Nature</i> 382:262-265, 1996.					
SE	AH	Bignon and Siminovitch, "Identification of PTP1C Mutation as the Genetic Defect in Motheaten and Viable Motheaten Mice: A Step Toward Defining the Roles of Protein Tyrosine Phosphatases in the Regulation of Hemopoietic Cell Differentiation and Function," <i>Clinical Immunology and Immunopathology</i> 73(2):168-179, 1994.					
SE	AI	Blair et al., "The Mouse Scurfy (sf) Mutation Is Tightly Linked to <i>Gatal</i> and <i>Tfe3</i> on the Proximal X Chromosome," <i>Mammalian Genome</i> 5:652-654, 1994.					
SE	AJ	Database Genbank Accession No. AI949471, August 1, 1998.					
SE	AK	Database Genbank Accession No. AJ005891, May 1, 1998.					
SE	AL	Derry et al., "The Mouse Homolog of the Wiskott-Aldrich Syndrome Protein (WASP) Gene Is Highly Conserved and Maps near the Scurfy (sf) Mutation on the X Chromosome," <i>Genomics</i> 29:471-477, 1995.					
SE	AM	Godfrey et al., "Fatal Lymphoreticular Disease in the Scurfy (sf) Mouse Requires T Cells that Mature in a sf Thymic Environment: Potential Model for Thymic Education," <i>Proc. Natl. Acad. Sci. USA</i> 88:5528-5532, 1991.					
SE	AN	Godfrey et al., "X-Linked Lymphoreticular Disease in the Scurfy (sf) Mutant Mouse," <i>American Journal of Pathology</i> 138(6):1379-1387, 1991.					
EXAMINER 				DATE CONSIDERED 5/28/02			
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### U.S. PATENT DOCUMENTS

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	BA						

### FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
							YES	NO
	BB							
	BC							
	BD							
	BE							
	BF							

### OTHER PRIOR ART *(Including Author, Title, Date, Pertinent Pages, Etc.)*

4-	BG	Lyon et al., "The Scurfy Mouse Mutant has Previously Unrecognized Hematological Abnormalities and Resembles Wiskott-Aldrich Syndrome," <i>Proc. Natl. Acad. Sci. USA</i> 87:2433-2437, 1990.
4-	BH	Rawlings et al., "Mutation of Unique Region of Bruton's Tyrosine Kinase in Immunodeficient XID Mice," <i>Science</i> 261:358-361, 1993.
4-	BI	Sugamura et al., "The Interleukin-2 Receptor $\gamma$ Chain: Its Role in the Multiple Cytokine Receptor Complexes and T Cell Development in XSCID," <i>Ann. Rev. Immuno.</i> 14:179-205, 1996.
4-	BJ	Veres et al., "The Molecular Basis of the Sparse Fur Mouse Mutation," <i>Science</i> 237:415-417, 1987.
	BK	
	BL	
	BM	
	BN	

EXAMINER

DATE CONSIDERED

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